Amendments to the Claims

1-43. (Cancelled)

44. (Currently amended) A metallocene-based ligand having a formula selected from the group consisting of Formula (I), Formula (II), Formula (III), Formula (IV), Formula (V), Formula (VI), Formula (VII), Formula (VIII), and Formula (IX):

wherein

W is phosphorus or arsenic;

M is a metal;

R¹ and R² are different from each other and are independently selected from the group consisting of unsubstituted branched-chain alkyl, unsubstituted straight-chain alkyl, unsubstituted alkoxy, unsubstituted alkylamino, unsubstituted cycloalkyl, unsubstituted cycloalkoxy, unsubstituted cycloalkylamino, unsubstituted carbocyclic aryl, unsubstituted carbocyclic aryloxy, unsubstituted heteroaryl, unsubstituted heteroaryloxy, unsubstituted carbocyclic arylamino, unsubstituted heteroarylamino, substituted branched-chain alkyl, substituted straight-chain alkyl, substituted alkoxy, substituted alkylamino, substituted cycloalkyl, substituted cycloalkoxy, substituted cycloalkylamino, substituted carbocyclic aryl, unsubstituted carbocyclic aryloxy, substituted heteroaryl, substituted heteroaryloxy, substituted carbocyclic arylamino, and substituted heteroarylamino;

R³ and R⁴ are independently selected from the group consisting of substituted branched-chain alkyl, substituted straight-chain alkyl, substituted cycloalkyl, substituted carbocyclic aryl, substituted heteroaryl, unsubstituted branched-chain alkyl, unsubstituted straight-chain alkyl, unsubstituted cycloalkyl, unsubstituted carbocyclic aryl, and unsubstituted heteroaryl;

n is an integer from 0 to 3;

m is an integer from 0 to 5;

Q is the group

wherein

R⁸ is selected from the group consisting of substituted straight-chain alky, substituted branched-chain alkyl, unsubstituted branched-chain alkyl, substituted cycloalkyl, unsubstituted cycloalkyl, substituted carbocyclic aryl, unsubstituted carbocyclic aryl, substituted heteroaryl, and unsubstituted heteroaryl; R⁹ and R^{10"} are independently selected from the group consisting of hydrogen, substituted straight-chain alkyl, substituted branched-chain alkyl, unsubstituted branched-chain alkyl, substituted cycloalkyl, unsubstituted carbocyclic aryl, unsubstituted carbocyclic aryl, unsubstituted heteroaryl, and unsubstituted heteroaryl; or

Q is selected from the group consisting of

$$R^{11}$$
 R^{12}
 R^{12}
 R^{12}

wherein

R⁶ and R⁷ are independently selected from the group consisting of substituted branched-chain alkyl, substituted straight-chain alkyl, substituted alkoxy, substituted alkylamino, substituted cycloalkyl, substituted cycloalkoxy, substituted cycloalkylamino, substituted carbocyclic aryl, substituted carbocyclic aryloxy, substituted heteroaryl, substituted heteroaryloxy, substituted carbocyclic arylamino, substituted heteroarylamino, unsubstituted branched-chain alkyl, unsubstituted straight-chain alkyl, unsubstituted alkoxy, unsubstituted alkylamino, unsubstituted cycloalkyl, unsubstituted cycloalkoxy, unsubstituted cycloalkylamino, unsubstituted carbocyclic aryl, unsubstituted carbocyclic aryloxy, unsubstituted heteroaryl, unsubstituted heteroaryloxy, unsubs

R⁸, R⁹, R¹⁰ and R¹⁰" are independently selected from the group consisting of hydrogen, substituted straight-chain alkyl, unsubstituted straight-chain alkyl, substituted branched-chain alkyl, unsubstituted branched-chain alkyl, substituted cycloalkyl, unsubstituted cycloalkyl, substituted carbocyclic aryl, unsubstituted carbocyclic aryl, substituted heteroaryl, and unsubstituted heteroaryl; R¹¹ is selected from the group consisting of OR¹³, SR¹³, NHR¹³, and NR¹³R¹⁴, wherein

R¹³ and R¹⁴ are independently selected from the group consisting substituted branched-chain alkyl, unsubstituted branched-chain alkyl, substituted cycloalkyl, unsubstituted cycloalkyl, substituted carbocyclic aryl, unsubstituted carbocyclic aryl, substituted heteroaryl, and

unsubstituted heteroaryl; R¹² is selected from the group consisting of hydrogen, halogen, OR¹³, SR¹³, NR¹³R¹⁴, substituted branched-chain alkyl, unsubstituted branched-chain alkyl, substituted cycloalkyl, unsubstituted carbocyclic aryl, unsubstituted carbocyclic aryl, substituted heteroaryl, and unsubstituted heteroaryl, and n' is 0 to 4;

R⁵ is selected from:

wherein R¹⁵, R¹⁶ and R¹⁷ are independently selected from the group consisting of hydrogen, haolgen halogen, OR¹³, SR¹³, NR¹³R¹⁴, substituted branched-chain alkyl, unsubstituted branched-chain alkyl, substituted cycloalkyl, unsubstituted cycloalkyl substituted carbocyclic aryl, unsubstituted carbocyclic aryl, substituted heteroaryl, and unsubstituted heteroaryl; and wherein the two geminal substituents R¹⁸ together are a doubly bonded oxygen atom, or each substituent R¹⁸ is individually hydrogen; and G is selected from the group consisting of -C(=O)NH-R*-NHCO-, -C(=O)-OR*O-C(=O)-, -C(=O)-R*C(=O)-, -CH=N-R*-N=CH-, -CH₂NH-R*-NHCH₂-, -CH₂NHC(=O)-R*-C(=O)NHCH₂-, -CH(R⁸)NH-R*-NH(CH(R⁸)-, -CH(R⁸)NHC(=O)-R*-C(=O)NHCH₂-, -CH₂NHC(=O)-, -C(=O)-NHCH₂-, -CH(R⁸)NH-R-NH(CH(R⁸)-, -CH(R⁸)NH-R-NH(CH(R⁸)-, -CH(R⁸)NH-R-NH(CH(R⁸)-, -CH(R⁸)NH-R-NH(CH(R⁸)-, -CH(R⁸)NH-R-NH(CH(R⁸)-, -CH(R⁸)NH-R-NH(CH(R⁸)-, -CH(R⁸)NHC(=O)-R-C(=O)NHCH(R⁸)-,

wherein R⁸ is, independently, as previously defined;

-R*- and -R- are selected from the group consisting of:

wherein R¹² is as previously defined;

wherein the two substituents R¹⁹ together are -(CH₂)_{m'}- or each substitutent R¹⁸ is independently selected from the group consisting of hydrogen, substituted branched-chain alkyl, unsubstituted branched-chain alkyl, substituted cycloalkyl, unsubstituted cycloalkyl, substituted carbocyclic

aryl, unsubstituted carbocyclic aryl, substituted heteroaryl, and unsubstituted heteroaryl; wherein the or each heteroatom is independently selected from sulphur, nitrogen, n' is an integer of from 0 to 4; and m' is an integer of from 1 to $\frac{8}{5}$.

with the proviso that the following compounds are excluded:

 $Fe(C_5H_5)(C_5H_3(CH_2NMe_2)(P-Ph-Me))$

Fe(C₅H₅)(C₅H₃(CHMeNMe₂)(P Ph Me)),

Fe(C₅H₅)(C₅H₃(CHMeNMe₂)(P Ph n-Bu)),

Fe(C₅H₅)(C₅H₃(CHMeNMe₂)(P Ph t-Bu)),

Fe(C₅H₅)(C₅H₃(CHMeNMe₂)(P n-Bu t-Bu)), and

Fe(C₅H₅)(C₅H₃(CHMeNMe₂)(P Ph-(3,5-bis(trifluoromethyl)phenyl))).

- 45. (Previously presented) The metallocene-based ligand of Claim 44, which is a diastereomer having Formula (IV), Formula (V), or Formula (VI).
- 46. (Previously presented) The metallocene-based ligand of Claim 44, which is an enantiomer having Formula (VII), Formula (VIII), or Formula (IX).
- 47. (Previously presented) The metallocene-based ligand of Claim 44, wherein the metallocene-based ligand is a phosphine or arsine having chirality at W, and wherein the metallocene-based ligand has at least one additional element of chirality selected from the group consisting of chirality at carbon, and axial chirality.
- 48. (Previously presented) The metallocene-based ligand of Claim 44, wherein the metallocene-based ligand is a diphosphine or diarsine having chirality at W, and wherein the metallocene-based ligand has two additional elements of chirality comprising chirality at carbon, and axial chirality.

- 49. (Previously presented) The metallocene-based ligand of Claim 44, wherein the metallocene is ferrocene.
- 50. (Previously presented) The metallocene-based ligand of Claim 44, wherein W is phosphorus.
- 51. (Previously presented) A catalyst or catalyst precursor in an asymmetric transformation reaction to generate a high enantiomeric excess of a formed compound, the catalyst or catalyst precursor comprising the metallocene-based ligand of Claim 44.
- 52. (Previously presented) A transition metal complex containing a transition metal coordinated to a ligand according to the metallocene-based ligand of Claim 44.
- 53. (Previously presented) A transition metal complex according to claim 52, wherein the transition metal is a Group VIb or a Group VIII metal.
- 54. (Previously presented) A method for preparing the metallocene-based ligand of Claim 44, comprising:

providing a metallocene-based substrate having a chiral directing substituent on one or both rings;

ortho-lithiating the metallocene-based substrate; and converting the ortho-lithiated metallocene-based substrate to obtain the metallocene-based ligand.

55. (Previously presented) The method according to Claim 54, wherein the metallocene-based ligand has Formula (I) or Formula (III), wherein the metallocene-based substrate has Formula (X'):

wherein R³ and R⁴ are independently selected from the group consisting of substituted branchedchain alkyl, substituted straight-chain alkyl, substituted cycloalkyl, substituted carbocyclic aryl, substituted heteroaryl, unsubstituted branched-chain alkyl, unsubstituted straight-chain alkyl, unsubstituted cycloalkyl, unsubstituted carbocyclic aryl, and unsubstituted heteroaryl;

and wherein X^* is a chiral directing group, wherein the step of converting the ortho-lithiated metallocene-based substrate comprises reacting the ortho-lithiated substrate with an R^1 substituted phosphine or arsine, and with an R^2 -bearing Grignard reagent or an R^2 -organolithium compound, then converting X^* to Q or G.

56. (Previously presented) A method according to Claim 55, wherein X* is selected from the group consisting of:

wherein

n is an integer from 0 to 3;

R^a and R^b are independently selected from the group consisting of substituted branched-chain alkyl, substituted straight-chain alkyl, substituted cycloalkyl, substituted carbocyclic aryl, substituted heteroaryl, unsubstituted branched-chain alkyl, unsubstituted straight-chain alkyl, unsubstituted cycloalkyl, unsubstituted carbocyclic aryl, and unsubstituted heteroaryl.

- 57. (Previously presented) The method according to claim 55, wherein the ortho-lithiation step is conducted using at least one lithiating agent selected from the group consisting of n-butyllithium, sec-butyllithium, and tert-butyllithium.
- 58. (Previously presented) The method according to claim 57, wherein the step of converting the ortho-lithiated metallocene-based substrate comprises reacting the ortho-lithiated metallocene-based substrate *in situ* with a dichlorophosphine of the formula R¹PCl₂ wherein R¹ is selected from the group consisting of unsubstituted branched-chain alkyl, unsubstituted straight-chain alkyl, unsubstituted alkoxy, unsubstituted alkylamino, unsubstituted cycloalkyl, unsubstituted cycloalkoxy, unsubstituted cycloalkylamino, unsubstituted carbocyclic aryl, unsubstituted carbocyclic aryloxy, unsubstituted heteroaryl, unsubstituted heteroaryloxy, unsubstituted carbocyclic arylamino, unsubstituted heteroarylamino, substituted branched-chain alkyl, substituted straight-chain alkyl, substituted alkoxy, substituted alkylamino, substituted cycloalkyl, substituted cycloalkoxy, substituted cycloalkylamino, substituted carbocyclic aryl, substituted carbocyclic aryloxy, substituted heteroaryl, substituted heteroaryloxy, substituted carbocyclic aryloxy, substituted heteroaryl, substituted heteroaryloxy, substituted carbocyclic aryloxy, substituted heteroaryl, substituted heteroaryloxy, substituted carbocyclic aryloxy, substituted heteroarylamino,

to yield an intermediate product, wherein the intermediate product is converted to obtain the metallocene-based ligand.

59. (Previously presented) The method according to Claim 58, further comprising reacting the intermediate product with an organometallic reagent of formula R²Z, wherein R² is selected from the group consisting of unsubstituted branched-chain alkyl, unsubstituted straight-chain alkyl, unsubstituted alkoxy, unsubstituted alkylamino, unsubstituted cycloalkyl, unsubstituted cycloalkylamino, unsubstituted carbocyclic aryl, unsubstituted

carbocyclic aryloxy, unsubstituted heteroaryl, unsubstituted heteroaryloxy, unsubstituted carbocyclic arylamino, unsubstituted heteroarylamino, substituted branched-chain alkyl, substituted straight-chain alkyl, substituted alkoxy, substituted alkylamino, substituted cycloalkyl, substituted cycloalkoxy, substituted cycloalkylamino, substituted carbocyclic aryl, substituted carbocyclic aryloxy, substituted heteroaryl, substituted heteroaryloxy, substituted carbocyclic arylamino, and substituted heteroarylamino;

wherein Z is Li or MgY, and wherein Y is a halide, to obtain a phosphorus chiral compound having formula (XI'):

$$R^{3} \qquad X^{*}$$

$$R^{2} \qquad R^{2}$$

$$R^{4} \qquad R^{1}$$

Formula (XI')

wherein the phosphorous chiral compound is converted to obtain the metallocene-based ligand.

- 60. (Previously presented) The method of Claim 59, wherein the metallocene-based ligand has Formula (I) or Formula (III).
- 61. (Previously presented) A method for preparing a metallocene-based ligand of Claim 44, comprising:

providing a compound of Formula (XXXVII):

wherein X is an achiral directing group;

subjecting the compound of Formula (XXXVII) to enantioselective mono-ortho-lithiation using

at least one lithiating agent selected from the group consisting of n-butyllithium, sec-butyllithium, and tert- butyllithium, wherein the mono-ortho-lithiation is conducted in the presence of a homochiral tertiary amine, whereby a chiral monolithium compound is obtained; reacting the chiral monolithium compound in situ with a dichlorophosphine of the formula R¹PCl₂ followed by reacting with an organometallic reagent of the formula R²Z, wherein R¹ and R² are different from each other and are independently selected from the group consisting of unsubstituted branched-chain alkyl, unsubstituted straight-chain alkyl, unsubstituted alkoxy, unsubstituted alkylamino, unsubstituted cycloalkyl, unsubstituted cycloalkoxy, unsubstituted cycloalkylamino, unsubstituted heteroaryl, unsubstituted heteroaryloxy, unsubstituted carbocyclic arylamino, unsubstituted heteroarylamino, substituted branched-chain alkyl, substituted straight-chain alkyl, substituted alkoxy, substituted alkylamino, substituted cycloalkyl, substituted cycloalkylamino, substituted cycloalkylamino, substituted cycloalkylamino, substituted cycloalkyl, substituted cycloalkylamino, substituted cheteroaryloxy, substituted carbocyclic aryloxy, substituted heteroaryl, substituted heteroaryloxy, substituted carbocyclic arylamino, and substituted heteroarylamino;

wherein Z is Li or MgY, and wherein Y is a halide, to obtain a phosphorus chiral compound having Formula (XXXVIII):

and converting the phosphorus chiral compound having Formula (XXXVIII) to the metallocene-based ligand, wherein the metallocene-based ligand has Formula (I) or Formula (III).

62. (Previously presented) The method according to Claim 61, wherein X is selected from the group consisting of:

$$NR^aR^b \neq SO_2R^a$$
 $NR^aR^b \neq P(O)R^aR^b$

wherein R^a and R^b are independently selected from the group consisting of substituted branched-chain alkyl, substituted straight-chain alkyl, substituted cycloalkyl, substituted carbocyclic aryl, substituted heteroaryl, unsubstituted branched-chain alkyl, unsubstituted straight-chain alkyl, unsubstituted cycloalkyl, unsubstituted carbocyclic aryl, and unsubstituted heteroaryl.

63. (Previously presented) A method for preparing a metallocene-based ligand of Claim 44, comprising:

providing a compound of the Formula (XXXIX):

wherein X* is a chiral directing group;

subjecting the compound of Formula (XXXIX) to bis-ortho-lithiation using at least one lithiating agent selected from the group consisting of n-butyllithium, sec-butyllithium, and tert-butyllithium, whereby a bislithium compound *in situ* with a dichlorophosphine of the formula R¹PCl₂ followed by reacting with an organometallic reagent of the formula R²Z wherein R¹ and R² are different from each other and are independently selected from the group consisting of unsubstituted branched-chain alkyl, unsubstituted straight-chain alkyl, unsubstituted alkoxy, unsubstituted alkylamino, unsubstituted cycloalkyl, unsubstituted cycloalkoxy, unsubstituted cycloalkylamino, unsubstituted heteroaryl, unsubstituted carbocyclic aryl, unsubstituted heteroarylamino, substituted branched-chain alkyl, substituted straight-chain alkyl, substituted alkoxy, substituted alkylamino, substituted cycloalkyl, substituted cycloalkoxy, substituted cycloalkylamino, substituted carbocyclic aryl, substituted cycloalkoxy, substituted heteroaryloxy, substituted carbocyclic aryloxy, substituted heteroaryloxy, substituted carbocyclic arylamino, and substituted heteroarylamino;

wherein Z is Li or MgY, and wherein Y is a halide, to obtain a phosphorus chiral compound having Formula (XXXX):

and converting the phosphorous chiral compound having Formula (XXXX) to the metallocene-based ligand, wherein the metallocene-based ligand has Formula (II).